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	Filing Date		2006-08-24
	First Named Inventor	Hirobumi Takahashi	
	Art Unit	1614	
	Examiner Name		
	Attorney Docket Number	BY0038P	

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	2	Chiou et al., "Nociceptin/Orphanin FQ Peptide Receptors: Pharmacology and Clinical Implications", Current Drug Targets, Vol. 8, pgs. 117-135, 2007.	<input type="checkbox"/>
	3	Niiyama et al., "6-Carboxy-5,7-Diarylcyclopentenol[1,2-b] Pyridine Derivatives: A Novel Class of Endothelin Receptor Antagonists", Biorg. Med. Chem., Vol. 10, pgs. 2461-2470, 2002.	<input type="checkbox"/>
	4	Pita et al., "A Simple, Efficient Method for Regioselective Synthesis of 7-Aminomehtyl-7,8-Dihydro-6H-Quinolin-5-ones, New Potential CNS Agents", Tetrahedron Letters, Vol. 41, pgs. 9829-9833, 2000.	<input type="checkbox"/>

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5	Ronzoni et al., "Lead Generation and Lead Optimisation Approches in the Discovery of Selective, Non-Peptide ORL-1 Receptor Agonists and Antagonists", Expert. Opin. Ther. Patents, Vol. 11, pgs. 525-546, 2001.	<input type="checkbox"/>
6	Zaratin et al., "Modification of Nociception and Morphine Tolerance by the Selective Opiate Receptor-Like Orphan Receptor Antagonist (-) Cis-1-Methyl-7[[4-2,6-Dichlorophenyl] Piperidin-1-yl]Methyl]-6,7,8,9-Tetrahydro-5H-Benzocyclophepten-5-ol (SB-612111)", J. of Pharmacology and Experimental Therapeutics, Vol. 308, pgs. 454-461, 2004.	<input type="checkbox"/>
7	Yoshizumi et al., "A Novel Class of Cycloalkano[b] Pyridines as Potent and Orally Active Opioid Receptor-Like 1 Antagonists with Minimal Binding Affinity to the hERG K+ Channel", J. Med. Chem, Vol. 51, pgs. 4021-4029, 2008.	<input type="checkbox"/>
8	Zaveri et al., "Peptide and NonPeptide Ligands for the Nociceptin/Orphanin FQ Receptor ORL1: Research Tools and Potential Therapeutic Agents", Life Sciences, Vol. 73, pgs. 663-678, 2003.	<input type="checkbox"/>

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